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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/776,419	02/10/2004	Shubh D. Sharma	70025-US04-404	2914
9629 7590 07/02/2007 MORGAN LEWIS & BOCKIUS LLP 1111 PENNSYLVANIA AVENUE NW WASHINGTON, DC 20004			EXAMINER SHIBUYA, MARK LANCE	
			ART UNIT 1639	PAPER NUMBER
			MAIL DATE 07/02/2007	DELIVERY MODE PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

<b>Office Action Summary</b>	<b>Application No.</b> 10/776,419	<b>Applicant(s)</b> SHARMA ET AL.	
	<b>Examiner</b> Mark L. Shibuya, Ph.D.	<b>Art Unit</b> 1639	

**-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --**

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 05 February 2007.
- 2a) ☐ This action is **FINAL**.                      2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 1-72 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☐ Claim(s) \_\_\_\_\_ is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☒ Claim(s) 1-72 are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All    b) ☐ Some    \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- \* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- |  |   |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892)   | 4) <input type="checkbox"/> Interview Summary (PTO-413)<br>Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)                       | 5) <input type="checkbox"/> Notice of Informal Patent Application                       |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)<br>Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____  |

### DETAILED ACTION

1. Claims 1-72 are pending.

#### ***Election/Restrictions***

2. The applicant is respectfully invited to note that claims 1-72 are listed as Groups "I, etc.", and "II, etc." and "III, etc.", but in actuality, contain within those claims a **large number of separate and distinct inventions**. Election of a single invention from within this group of claims is required as specifically set forth (see Further Restriction, below).

3. Restriction to one of the following inventions is required under 35 U.S.C. 121:

I, etc.        Claims 1-33, 38-49 drawn to methods of deriving a peptidomimetic comprising selecting a biologically active metallopeptide, modeling a non-peptidic ring structure that is superimposable on a template space, and forming a peptidomimetic, classifiable in class 530, subclass 323.

II, etc.        Claims 34, 35, 36, 37, drawn to peptidomimetic, classifiable in class 514, subclass 2.

III, etc.        Claims 50-72, drawn to a method of deriving a peptidomimetic comprising, designing and constructing a library of amino acid sequences, complexing each library member to a metal ion to forming a library of metallopeptides, and screening the library of metallopeptides, selecting a

metallopeptide, modeling a non-peptidic ring structure that is superimposable on a template space, and forming a peptidomimetic classifiable in class 435, subclass DIG 15.

#### Further Restriction

- A) In addition each of Groups I, etc., II, etc., and III, etc., (methods and compounds comprising peptidomimetics, such as the peptidomimetics in claims 33 and 35) reads on patentably distinct Groups. Groups I, etc, II, etc., and III, etc., are further divided into multiple groups each representing a different molecular core ring structure. The compounds within each Group comprise a different core ring structure that has no unifying structural relationship with the other ring core structures, which results in a shared functional property among the plurality of core ring structures. Thus a further restriction is applied to each of Groups I, etc., II, etc., and III, etc..

The inventions are distinct, each from the other because of the following reasons:

The claims of Group I, etc., II, etc., and III, etc., are drawn to compounds with different ring structures.

If one of Groups I, etc., or II, etc., is elected, the elected **further restricted Group** *must* result in a single specific generic compound, *i.e.*, a

product comprising a single molecular core ring structure. If an Invention of Groups I, etc., II, etc., or III, etc., is elected, a specific molecular structure must be elected as a species, (see below requirement for election of species).

*For this response to be complete, applicants should provide the structure of the elected core ring structure and list all of the claims readable upon the elected core.*

This requirement is not to be taken as an election of species, but rather as an election of a single invention, since each product is assumed to be a patentably distinct invention, absent evidence to the contrary.

The Inventions of Groups I, etc. and III, etc and the invention of Group II, etc., are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make another and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case the peptidomimetic product of Group II, etc., may be generated using X-ray crystallography or NMR, which is different method from methods comprising modeling a non-peptidic ring structure that is superimposable on a template space and adding elements to occupy similar descriptor spaces, as in Groups I, etc., and III, etc.

Inventions of Groups I, etc., and III, etc., are directed to related processes. The related inventions are distinct if the (1) the inventions as claimed are either not capable of use together or can have a materially different design, mode of operation, function, or effect; (2) the inventions do not overlap in scope, i.e., are mutually exclusive; and (3) the inventions as claimed are not obvious variants. See MPEP § 806.05(j). In the instant case, the inventions as claimed have materially different design and modes of operation and function, because the method of Group III, etc., comprising the design, formation, and screening of two different libraries and so is materially different from the method of Group I, etc. Furthermore, the inventions as claimed do not encompass overlapping subject matter and there is nothing of record to show them to be obvious variants.

Because these inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required because the inventions have acquired a separate status in the art in view of their different classification, restriction for examination purposes as indicated is proper.

Because these inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required because the inventions require a different field of search (see MPEP § 808.02), restriction for examination purposes as indicated is proper.

Because these inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required because the inventions have acquired a separate status in the art due to their

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recognized divergent subject matter, restriction for examination purposes as indicated is proper.

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. **Process claims that depend from or otherwise include all the limitations of the patentable product** will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai*, *In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. **Failure to do so may result in a loss of the right to rejoinder.**

Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

#### *Requirement for Election of Species*

4. Claims 1-72 are generic to the following disclosed patentably distinct species:

Applicant must elect an ultimate species of peptidomimetic, defined as to atom and

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bond. The species are independent or distinct because the different peptidomimetic have different structures that provide materially different function and effect. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

5. Claims 1-72 are generic to the following disclosed patentably distinct species: a species of metallopeptide, including specificity of the biological activity of the metallopeptide, ( as in claims 25-26 and 27-30). The species are independent or distinct because the different metallopeptides have different structures that provide materially different function and effect. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

6. Claims 1-72 are generic to the following disclosed patentably distinct species: Methods comprising a species of template space. The species are independent or distinct because the different template spaces have materially different design, modes of operation, function and effect. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

7. Claims 1-72 are generic to the following disclosed patentably distinct species: Methods comprising a species of descriptor space. The species are independent or distinct because the different descriptor spaces have materially different design, modes



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of operation, function and effect. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

8. Claims 1-72 generic to the following disclosed patentably distinct species: A target of interest. The species are independent or distinct because the different targets of interest have materially different function and effect. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

9. This application contains claims directed to the following patentably distinct species: A species of at least four atoms: Claim 4:  $N_3S_1$ ; claim 5:  $N_2S_2$ . The species are independent or distinct because the different the different at least four atoms have different molecular structures that confer materially different design.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1, 3 are generic.

10. This application contains claims directed to the following patentably distinct species: Claims 13 and 14: A method comprising species of template space that are defined by one ring or two rings of a tricyclic ring structure. The species are independent or distinct because the different species of template space by materially different design, modes of operation, function and effect.

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Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1, 12 are generic.

11. This application contains claims directed to the following patentably distinct species: Methods comprising species of comparing the biological activity of the peptidomimetic to the metallopeptide: Claims 19, 20, 21, 22, 23, 24, 42-48, 51-52. The species are independent or distinct because the different species of comparison have materially different modes of operation, function and effect.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1, 2, 38, 40, 50, are generic.

12. This application contains claims directed to the following patentably distinct species: A method comprising a library of amino acid sequences comprising species of residues: Claims 54-57. The species are independent or distinct because the different residues have materially different design and modes of operation.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claim 50 is generic.

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13. Claim 50 is generic to the following disclosed patentably distinct species: A species of a value of  $n$ . The species are independent or distinct because the different values of  $n$  have materially different effect, particularly in the step of dividing into primary structures, (see claim 71). Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

14. This application contains claims directed to the following patentably distinct species: Methods comprising species of cysteine residue substitutions: claims 63-66. The species are independent or distinct because the different species of cysteine residue substitutions result in methods of materially different design.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claim 50, 63 are generic.

15. Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which depend from or otherwise require all the limitations

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of an allowable generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species.

MPEP § 809.02(a).

16. Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Should applicant traverse on the ground that the inventions or species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the inventions or species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103(a) of the other invention.

17. For this response to be complete and for search purposes, applicants should provide the ***chemical structure of elected compounds species***, wherein the specific formula substituents of the above identified elected species is defined, either by picture

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or by expressing the species in terms of the variables of the formula. Thus, applicant should provide, for search purposes, a chemical structure of a particular elected distinct species claimed, defined as to atom and bond; and a molecular core ring structure for the Invention of the Group elected, as required above in the instant Requirement for Restriction/Election. The provided chemical structure of the elected species must depict a single molecule, from which a search is to commence.

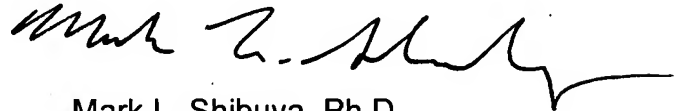
18. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

19. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Mark Shibuya, whose telephone number is (571) 272-0806. The examiner can normally be reached on M-F, 8:30AM-5:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Dr. J. Douglas Schultz can be reached on (571) 272-0763. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.



Mark L. Shibuya, Ph.D.  
Primary Examiner  
Art Unit 1639